Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	2	"5496872".did.	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 15:28
S1	1757	530/324.ccls. and coupling	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:39
S2	1334	530/324.ccls. and coupling SAME peptide	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:40
S3	0	530/324.ccls. and coupling SAME peptide and "NH-CO-A-CO-S"	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:40
S4	0	"NH-CO-A-CO-S"	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:40
S5	0	"NH-CO-A-CO"	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:40
S6	3	530/324.ccls. and coupling SAME peptide and alkylene and arylene	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:41
S7	247	coupling SAME peptide and alkylene and arylene	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:41
S8	202	coupling SAME peptide and alkylene SAME arylene	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 12:41

S9	3	coupling SAME peptide SAME alkylene SAME arylene	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/05/13 15:25

- Search History will be lost after eight hours of inactivity.
- To combine searches use # before search number, e.g., #2 AND #6.
- Search numbers may not be continuous; all searches are represented.
- Click on query # to add to strategy

Search	Most Recent Queries	Time	Result
#4	Search **bradykinin and dimers**	15:07:21	15
#3	Search **stewart and bradykinin and dimers**	15:07:10	4
#2	Search **stewart and bradykinin and 1999**	14:53:11	4
#1	Search **stewart and bradykinin**	14:53:05	173

Clear History

E-Mail Notice - SEARCH

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=> d his full

     (FILE 'HOME' ENTERED AT 13:11:45 ON 13 MAY 2005)

     FILE 'HCAPLUS' ENTERED AT 13:11:51 ON 13 MAY 2005
L1             1 SEA ABB=ON  PLU=ON  (GB2000-12083# OR GB1999-20397# OR
                 WO2000-GB3306#)/AP,PRN

     FILE 'REGISTRY' ENTERED AT 13:13:58 ON 13 MAY 2005

     FILE 'HCAPLUS' ENTERED AT 13:14:01 ON 13 MAY 2005
L2             TRA L1 1- RN :       47 TERMS

     FILE 'REGISTRY' ENTERED AT 13:14:01 ON 13 MAY 2005
L3            47 SEA ABB=ON  PLU=ON  L2
L4            26 SEA ABB=ON  PLU=ON  L3 AND S/ELS
L5            19 SEA ABB=ON  PLU=ON  L4 AND S=1
L6             1 SEA ABB=ON  PLU=ON  L5 AND C59H83N12O37P5S
L7               STR
L8             0 SEA CSS SAM L7
L9               SCR 2039 OR 2041 OR 2050 OR 2049 OR 2048 OR 2053 OR 2052 OR 204
L10              SCR 2007 AND 2021 AND 1993
L11            0 SEA CSS SAM L7 AND L10 NOT L9
L12            0 SEA CSS FUL L7 AND L10 NOT L9
L13            0 SEA SSS SAM L7 AND L10 NOT L9
L14           14 SEA SSS FUL L7 AND L10 NOT L9

     FILE 'HCAPLUS' ENTERED AT 13:47:35 ON 13 MAY 2005

     FILE 'REGISTRY' ENTERED AT 13:47:39 ON 13 MAY 2005
                SAV TEM AUD222F2/A L14

     FILE 'HCAPLUS' ENTERED AT 13:47:54 ON 13 MAY 2005
L15            5 SEA ABB=ON  PLU=ON  L14

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L16            2 SEA ABB=ON  PLU=ON  L14
                 SEL AN
                 EDIT E1-E2 /AN /OREF

     FILE 'HCAPLUS' ENTERED AT 13:48:29 ON 13 MAY 2005
L17            2 SEA ABB=ON  PLU=ON  ("CA53:12186A"/OREF OR "CA56:11449G"/OREF)

L18            7 SEA ABB=ON  PLU=ON  L15 OR L17

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  Please note that search-term pricing does apply when
  conducting SmartSELECT searches.

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Search done by Noble Jarrell

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* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added,    *
* effective March 20, 2005.  A new display format, IDERL, is now     *
* available and contains the CA role and document type information.  *
*                                                                    *
**********************************************************************

Crossover limits have been increased.  See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more
information enter HELP PROP at an arrow prompt in the file or refer
to the file summary sheet on the web at:
http://www.cas.org/ONLINE/DBSS/registryss.html

=> d que sta l14
L7                STR
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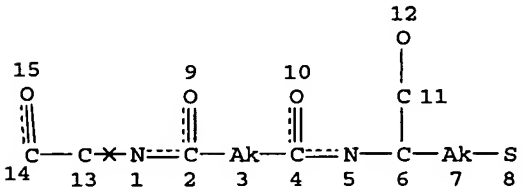


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CONNECT IS M1  RC AT   1
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GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
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STEREO ATTRIBUTES: NONE
L9                SCR 2039 OR 2041 OR 2050 OR 2049 OR 2048 OR 2053 OR 2052 O
R 2043 OR 2054
L10               SCR 2007 AND 2021 AND 1993
L14           14 SEA FILE=REGISTRY SSS FUL L7 AND L10 NOT L9

100.0% PROCESSED 350225 ITERATIONS                          14 ANSWERS
SEARCH TIME: 00.00.18


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FILE LAST UPDATED: 12 May 2005 (20050512/ED)

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=> d all hitstr l18 tot

L18 ANSWER 1 OF 7 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:66410 HCAPLUS
DN 138:394839
ED Entered STN: 28 Jan 2003
TI Zinc complexation of glutathione and glutathione-derived peptides
AU Gelinsky, M.; Vogler, R.; Vahrenkamp, H.
CS Institut fur Anorganische und Analytische Chemie der Universitat Freiburg, Freiburg, D-79104, Germany
SO Inorganica Chimica Acta (2003), 344, 230-238
CODEN: ICHAA3; ISSN: 0020-1693
PB Elsevier Science B.V.
DT Journal
LA English
CC 78-7 (Inorganic Chemicals and Reactions)
Section cross-reference(s): 34
OS CASREACT 138:394839
AB Glutathione (γ-Glu-Cys-Gly, GSH, 1) forms a binary Zn complex L2Zn3 (1.1) which is polymeric. Its triply blocked and purely S-functional form [4-NO2-Bz]-γ-Glu(Cys-Gly-OEt)-OEt (2) yields a polymeric complex L2Zn (2.1) and a monomeric pyrazolylborate Zn complex TpZn-SR (2.2). Doubly O-protected GSH was converted with histidine and cysteine to the difunctional tetrapeptides NAc-His-γ-Glu(Cys-Gly-OEt)-OEt (3) and NAc-Cys-γ-Glu(Cys-Gly-OEt)-OEt (4). Peptide 3 could be converted to the oligomeric Zn halide complex L·ZnCl (3.1). The Zn complexation of peptide 4 was studied by potentiometric titrns., revealing that the dominating species in solution are [ZnL(LH)]- and [ZnL2]2-, both with a ZnS4 coordination. In contrast, the isolated complex is the polymeric species LZn (4.1). With pyrazolylborate Zn units a monomeric dizinc complex TpZn-S.apprx..apprx.S-ZnTp (4.2) was obtained. N-protected GSH was extended by two histidine or cysteine units to the pentapeptides H-γ-Glu(Cys-Gly-His-OMe)-His-OMe (5) and H-γ-Glu(Cys-Gly-Cys-OMe)-Cys-OMe (6). Compds. 5 and 6 formed ill-defined polymeric Zn complexes. For 5, [L·Zn]ClO4, and for 6, [L3Zn5]CF3COO could be obtained anal. pure.
ST glutathione deriv peptide prepn complexation zinc; zinc complex glutathione deriv peptide prepn
IT Peptides, preparation
RL: SPN (Synthetic preparation); PREP (Preparation)
(zinc complexes; preparation of zinc(II) complexes of glutathione and of glutathione-derived peptide derivs.)
IT 183113-39-5
RL: RCT (Reactant); RACT (Reactant or reagent)
(complexation with glutathione-derived peptides)
IT 70-18-8, Glutathione, reactions 1499-46-3 27486-84-6 27486-87-9 183498-47-7 302325-91-3 528530-58-7
RL: RCT (Reactant); RACT (Reactant or reagent)
(for preparation of glutathione-derived peptides and their zinc(II) complexes)
IT 528530-74-7P 528530-77-0P 528530-80-5P 528530-82-7P 528530-84-9P 528530-86-1P 528530-88-3P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(for preparation of glutathione-derived peptides and their zinc(II) complexes)
IT 528530-61-2P 528530-64-5P 528530-66-7P 528530-69-0P 528530-72-5P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT

(Reactant or reagent)
(preparation and complexation with zinc(II))
IT 7440-66-6DP, Zinc, complexes of glutathione-derived peptides
35436-84-1DP, zinc complex 528530-63-4DP, zinc complex 528530-68-9DP, zinc chloro complex **528530-71-4DP**, zinc complex 528530-90-7P
528530-92-9P 528530-96-3P 528530-98-5P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
IT 122-04-3, 4-Nitrobenzoyl chloride
RL: RCT (Reactant); RACT (Reactant or reagent)
(protecting group for preparation of glutathione-derived peptides and their zinc(II) complexes)
RE.CNT 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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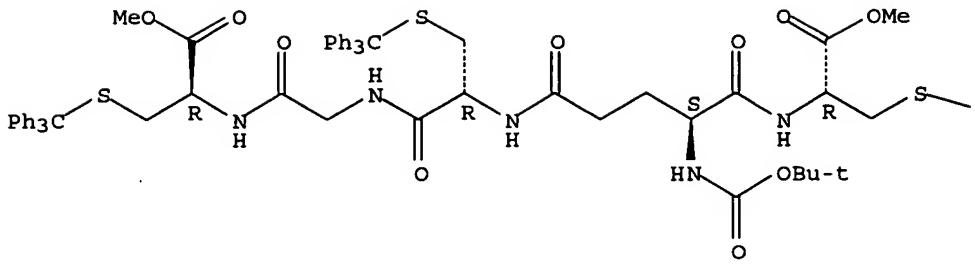
(26) Vallee, B; Acc Chem Res 1993, V26, P543 HCAPLUS
(27) Vogler, R; Inorg Chim Acta 2002, V329, P1
IT **528530-88-3P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(for preparation of glutathione-derived peptides and their zinc(II) complexes)
RN 528530-88-3 HCAPLUS
CN L-Cysteine, S-(triphenylmethyl)-L-cysteinylglycyl-S-(triphenylmethyl)-, methyl ester, (1'→1)-amide with N-[(1,1-dimethylethoxy)carbonyl]-L-α-glutamyl-S-(triphenylmethyl)-L-cysteine 2'-methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



PAGE 1-B

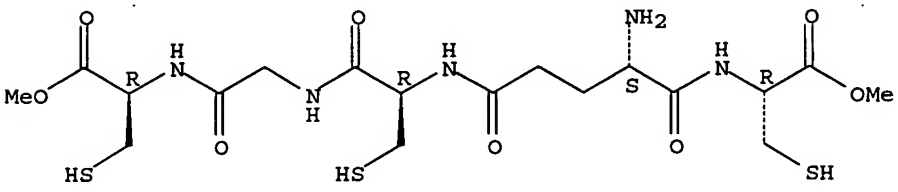
—CPh3

IT 528530-72-5P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and complexation with zinc(II))
RN 528530-72-5 HCAPLUS
CN L-Cysteine, L-cysteinylglycyl-, methyl ester, (1'→1)-amide with L-α-glutamyl-L-cysteine 2'-methyl ester, mono(trifluoroacetate) (salt) (9CI) (CA INDEX NAME)

CM 1

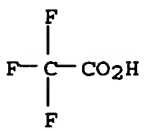
CRN 528530-71-4
CMF C18 H31 N5 O8 S3

Absolute stereochemistry.



CM 2

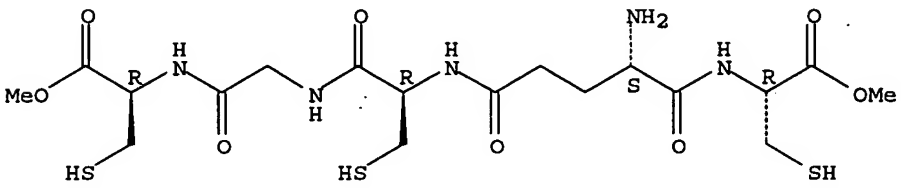
CRN 76-05-1
CMF C2 H F3 O2



IT 528530-71-4DP, zinc complex
RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)
RN 528530-71-4 HCAPLUS
CN L-Cysteine, L-cysteinylglycyl-, methyl ester, (1'→1)-amide with L-α-glutamyl-L-cysteine 2'-methyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L18 ANSWER 2 OF 7 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 2002:866867 HCAPLUS
DN 137:370487
ED Entered STN: 15 Nov 2002
TI Manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides
IN Kudo, Hiroto; Endo, Takeshi
PA JSR Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 4 pp.
CODEN: JKXXAF
DT Patent
LA Japanese
IC ICM C08G075-14
CC 35-5 (Chemistry of Synthetic High Polymers)
Section cross-reference(s): 28
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 2002327061	A2	20021115	JP 2001-132057	20010427
PRAI JP 2001-132057		20010427		

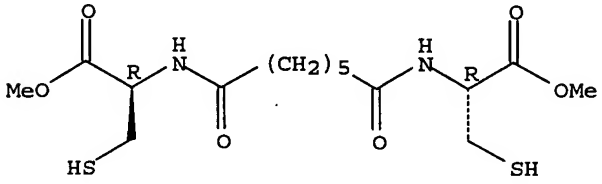
CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
JP 2002327061	ICM	C08G075-14

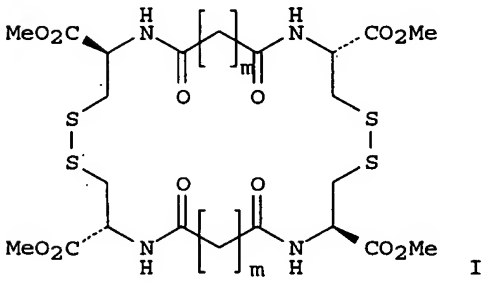
AB Polydisulfides are manufactured by ring-opening polymerization of cyclic disulfides obtained from dithiols having cysteine groups at the both terminals. Thus, L-Cys Me ester HCl salt was amidated with pimeloyl dichloride, dimerized, and polymerized at 100° for 24 h in DMF to give 90% a polydisulfide with average mol. weight 25,000.
ST cyclic cysteine ring opening polymn; dithiol cyclic ring opening polymn; polydisulfide manuf ring opening polymn
IT Polysulfides
RL: IMF (Industrial manufacture); PREP (Preparation)
(polyamide-; manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides)
IT Polyamides, preparation
RL: IMF (Industrial manufacture); PREP (Preparation)
(polysulfide-; manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides)
IT Polymerization
(ring-opening; manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides)
IT 474511-29-0P 474511-30-3P
RL: IMF (Industrial manufacture); PREP (Preparation)
(manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides)
IT 256953-91-0P 400652-92-8P
RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
(manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides)

IT 142-79-0, Pimeloyl dichloride 18598-63-5, L-Cysteine Methyl ester hydrochloride
RL: RCT (Reactant); RACT (Reactant or reagent)
(manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides)
IT 400652-92-8P
RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
(manufacture of polydisulfides by ring-opening polymerization of cyclic disulfides)
RN 400652-92-8 HCAPLUS
CN L-Cysteine, N,N'-(1,7-dioxo-1,7-heptanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



L18 ANSWER 3 OF 7 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 2001:749822 HCAPLUS
DN 136:200456
ED Entered STN: 15 Oct 2001
TI Efficient synthesis of macrocycles by oxidation of cysteine-based dithiols
AU Kudo, H.; Sanda, F.; Endo, T.
CS Chemical Resources Laboratory, Tokyo Institute of Technology, Midori-ku, Yokohama, 226-8503, Japan
SO Tetrahedron Letters (2001), 42(44), 7847-7850
CODEN: TELEAY; ISSN: 0040-4039
PB Elsevier Science Ltd.
DT Journal
LA English
CC 34-3 (Amino Acids, Peptides, and Proteins)
Section cross-reference(s): 28
OS CASREACT 136:200456
GI



I

AB The cysteine-bridged macrocycles I (m = 1-6), 22-32-membered cyclic dimers could be synthesized with a high reagent concentration (1 M) in excellent yields by the oxidation of the cysteine-based compds. having dithiol groups.
ST cysteine methyl ester amidation alkanedicarboxyl chloride; dithiol cysteine based prepn dimerization polymn; macrocyclic amide polymethylene bridged prepn; polymer cysteine bridged prepn
IT Acid halides
RL: RCT (Reactant); RACT (Reactant or reagent)
(chlorides, diacid; preparation of cysteine-bridged macrocycles via oxidation

of dithiols)
IT Macrocyclic compounds
RL: SPN (Synthetic preparation); PREP (Preparation)
(cysteine-bridged, amides; preparation by oxidation of cysteine-based dithiols)
IT Thiols (organic), preparation
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(dithiols, cysteine-based; preparation of cysteine-bridged macrocycles by oxidation of cysteine-based dithiols)
IT Oxidation
(of cysteine-based dithiols to cysteine-bridged macrocycles)
IT 111-50-2, Hexanedioyl dichloride 142-79-0, Heptanedioyl dichloride 543-20-4, Succinoyl chloride 1663-67-8, Malonyl chloride 2485-62-3, L-Cysteine methyl ester 2873-74-7, Pentanedioyl dichloride 10027-07-3, Octanedioyl dichloride
RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of cysteine-based dithiols for oxidation to cysteine-bridged macrocycles)
IT 400652-95-1P 400652-96-2P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of cysteine-based polymers)
IT 400652-88-2
RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent)
(preparation of cysteine-bridged macrocycles by oxidation of dithiols)
IT 400652-89-3P 400652-90-6P 400652-91-7P 400652-92-8P 400652-93-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of cysteine-bridged macrocycles by oxidation of dithiols)
IT 256953-80-7P 256953-88-5P 256953-91-0P 256953-95-4P 400652-94-0P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of cysteine-bridged macrocycles by oxidation of dithiols)
RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
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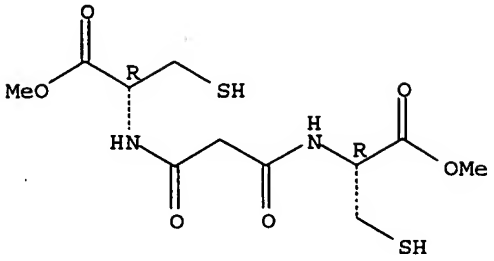
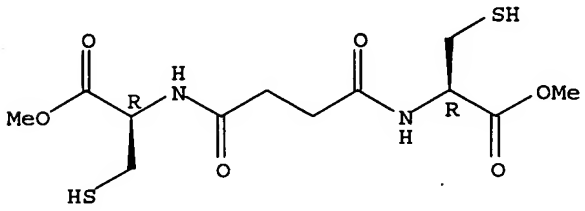
(23) Sprengard, U; Angew Chem, Int Ed Engl 1996, V35, P321 HCAPLUS
(24) Zhang, I; J Am Chem Soc 1997, V119, P2363
IT 400652-88-2
RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent)
(preparation of cysteine-bridged macrocycles by oxidation of dithiols)
RN 400652-88-2 HCAPLUS
CN L-Cysteine, N,N'-(1,3-dioxo-1,3-propanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



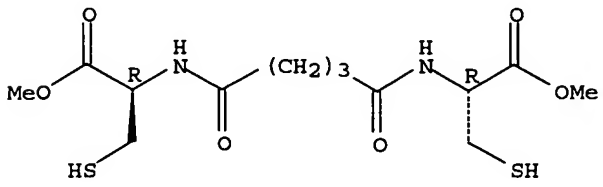
IT 400652-89-3P 400652-90-6P 400652-91-7P
400652-92-8P 400652-93-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of cysteine-bridged macrocycles by oxidation of dithiols)
RN 400652-89-3 HCAPLUS
CN L-Cysteine, N,N'-(1,4-dioxo-1,4-butanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



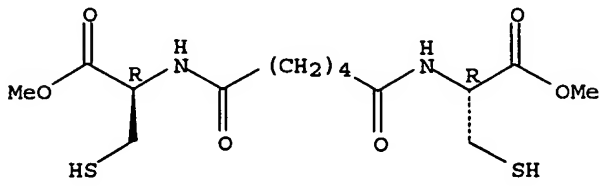
RN 400652-90-6 HCAPLUS
CN L-Cysteine, N,N'-(1,5-dioxo-1,5-pentanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



RN 400652-91-7 HCAPLUS
CN L-Cysteine, N,N'-(1,6-dioxo-1,6-hexanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).

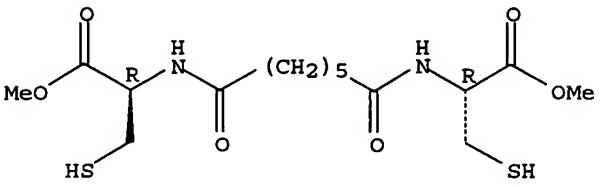


RN 400652-92-8 HCAPLUS

Search done by Noble Jarrell

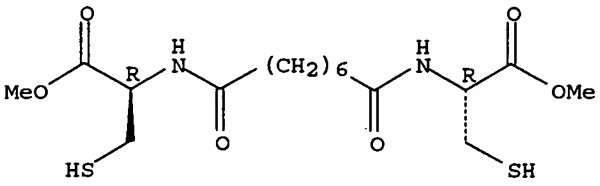
CN L-Cysteine, N,N'-(1,7-dioxo-1,7-heptanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



RN 400652-93-9 HCAPLUS
CN L-Cysteine, N,N'-(1,8-dioxo-1,8-octanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (+).



L18 ANSWER 4 OF 7 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 1976:165204 HCAPLUS
DN 84:165204
ED Entered STN: 12 May 1984
TI Insulin, insulin analogs, and derivatives
IN Geiger, Rolf; Obermeier, Rainer
PA Hoechst A.-G., Fed. Rep. Ger.
SO Ger. Offen., 14 pp.
CODEN: GWXXBX
DT Patent
LA German
IC C07C; A61K
CC 34-4 (Synthesis of Amino Acids, Peptides, and Proteins)
FAN.CNT 1

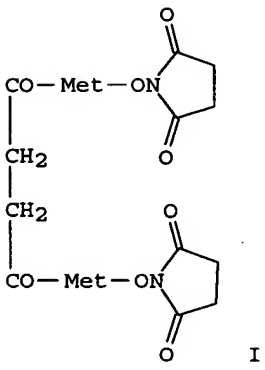
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2428412	A1	19760115	DE 1974-2428412	19740612
	NL 7506754	A	19751216	NL 1975-6754	19750606
	CH 613185	A	19790914	CH 1975-7416	19750609
	US 4014861	A	19770329	US 1975-585604	19750610
	CA 1044227	A1	19781212	CA 1975-229026	19750610
	DK 7502631	A	19751213	DK 1975-2631	19750611
	SE 7506695	A	19751215	SE 1975-6695	19750611
	AU 7582038	A1	19761216	AU 1975-82038	19750611
	BE 830186	A1	19751212	BE 1975-157293	19750612
	FR 2274605	A1	19760109	FR 1975-18372	19750612
	JP 51023290	A2	19760224	JP 1975-73026	19750612
	JP 58011427	B4	19830302		
	US 32015	E	19851029	US 1979-22852	19790322
PRAI	DE 1974-2428412	A	19740612		
	US 1975-585604	A5	19750610		

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
DE 2428412	IC	C07CIC A61K
US 4014861	NCL	530/303.000; 530/409.000; 930/010.000; 930/DIG.621

US 32015 NCL 530/303.000; 530/339.000; 930/010.000; 930/260.000; 930/DIG.621

GI



I

AB Blocking the 1st residue of insulin A chain tetrasulfonate with the methionine derivative I followed by reaction with NB1-(trifluoroacetyl)insulin B chain disulfonate in 4-ethylmorpholine at pH 8-11 gave a methionine bridged insulin derivative which then underwent disulfide bridge formation in thioglycol under N followed by cleavage of the blocking groups and the methionine bridging group to give 38% insulin with biol. activity.

ST insulin methionine bridge

IT 59113-60-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and deblocking of)

IT 59006-17-6P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and deesterification of)

IT 59006-18-7P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and reaction with hydroxysuccinimide)

IT 59006-19-8P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction with insulin A chain tetrasulfonate)

IT 59217-92-4P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction with insulin A chain tetrasulfonate methioine derivs.)

IT 18152-38-0P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction with methionine derivs.)

IT 11137-90-9P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction with trifluoroacetic acid methyl ester)

IT 59006-20-1P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

IT 431-47-0
RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with insulin derivs.)

IT 6066-82-6

RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with methionine derivs.)

IT 108-30-5
RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with methionine methyl ester)

IT 2491-18-1
RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with succinic anhydride)

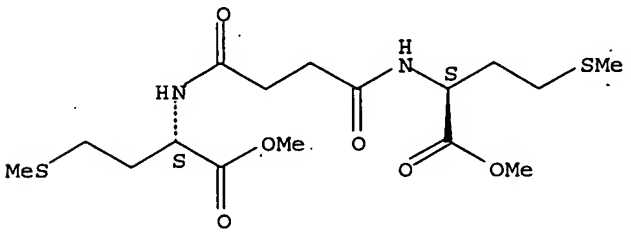
IT 11070-73-8
RL: RCT (Reactant); RACT (Reactant or reagent)
(using methionine bridging groups)

IT **59006-17-6P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and deesterification of)

RN 59006-17-6 HCAPLUS

CN L-Methionine, N,N'-(1,4-dioxo-1,4-butanediyl)bis-, dimethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

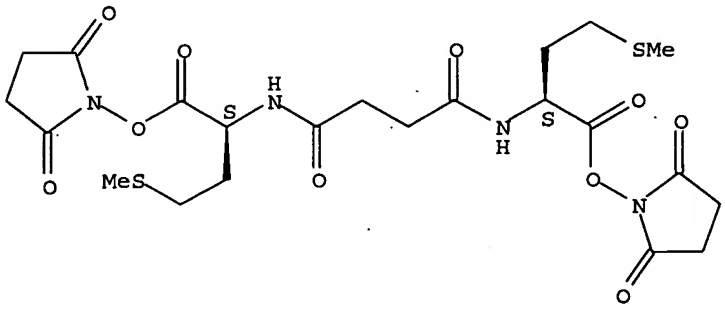


IT **59006-19-8P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction with insulin A chain tetrasulfonate)

RN 59006-19-8 HCAPLUS

CN Butanediamide, N,N'-bis[1-[[(2,5-dioxo-1-pyrrolidinyl)oxy]carbonyl]-3-(methylthio)propyl]-, [S-(R*,R*)]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

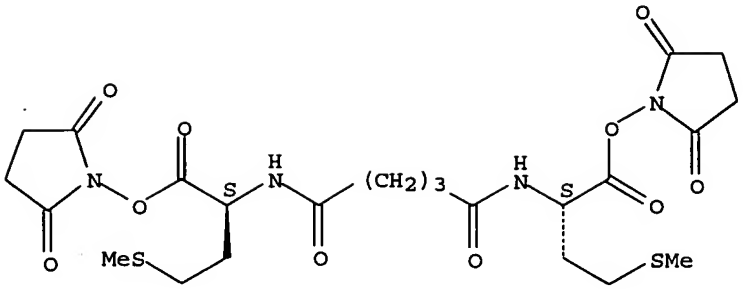


IT **59006-20-1P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 59006-20-1 HCAPLUS

CN Pentanediamide, N,N'-bis[1-[[(2,5-dioxo-1-pyrrolidinyl)oxy]carbonyl]-3-(methylthio)propyl]-, [S-(R*,R*)]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L18 ANSWER 5 OF 7 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 1973:537505 HCAPLUS
DN 79:137505
ED Entered STN: 12 May 1984
TI Sulfur-containing amino acid derivatives useful in therapeutics and in cosmetics
IN Dussourd, Lucien; Cousse, Henri; Bonnaud, Bernard
PA Fabre, Pierre, S. A.
SO Fr. Demande, 8 pp. Div. of Fr. 2,159,183.
CODEN: FRXXBL
DT Patent
LA French
IC A61K; C07C; C07D
CC 34-2 (Synthesis of Amino Acids, Peptides, and Proteins)
Section cross-reference(s): 1, 62

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI FR 2159183	A1	19730622	FR 1971-40215	19711109
FR 2159183	B1	19750606		
PRAI FR 1971-40215	A	19711109		

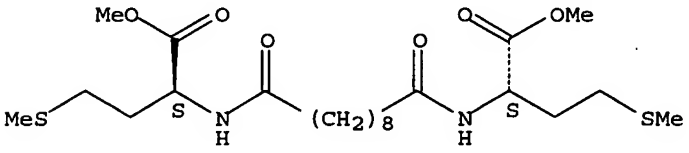
CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
FR 2159183	IC	A61KIC C07CIC C07D

AB MeSCH2CH2CH(COR)NHCO(CH2)8CONHCH(COR)CH2CH2SMe (I, R = OH, ONa, OMe, NH2) and related compds. were prepared for treating acne and seborrhea. Thus I (R = OH) was obtained by treating methionine with sebacoyl chloride.
ST methionine sebacoyl acne seborrhea
IT Acne
(bis(thiomethyl)piperazinedione for treatment of)
IT Seborrhea
(pyridoxine N-sebacoylmethioninate for treatment of)
IT Shampoos
(sodium N-sebacoylmethioninate containing)
IT 49761-96-8
RL: RCT (Reactant); RACT (Reactant or reagent)
(acne treatment composition containing)
IT 49761-97-9 50794-98-4
RL: RCT (Reactant); RACT (Reactant or reagent)
(antiseborrheal composition containing)
IT 49761-90-2P 49761-91-3P 49761-92-4P 49761-94-6P
49761-95-7P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
IT 111-19-3
RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with methionine methyl ester)
IT 10332-17-9 19298-72-7
RL: RCT (Reactant); RACT (Reactant or reagent)
(reaction of, with sebacoyl chloride)

IT 111-20-6, reactions
RL: RCT (Reactant); RACT (Reactant or reagent)
(with methioninamide)
IT 63-68-3, reactions
RL: RCT (Reactant); RACT (Reactant or reagent)
(with sebacoyl chloride)
IT **49761-92-4P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 49761-92-4 HCAPLUS
CN L-Methionine, N,N'-(1,10-dioxo-1,10-decanediyl)bis-, dimethyl ester (9CI)
(CA INDEX NAME)

Absolute stereochemistry.



L18 ANSWER 6 OF 7 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 1962:60315 HCAPLUS
DN 56:60315
OREF 56:11449f-h
ED Entered STN: 22 Apr 2001
TI Mucic acid derivatives
IN Morel, Charles J.
PA J. R. Geigy A.-G.
DT Patent
LA Unavailable
INCL 12P
CC 27 (Aliphatic Compounds)

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CH 354788		19610731	CH	19570415

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
CH 354788	INCL	12P

AB Compds. with strong antiphlogistic activity were prepared by treating mucic acid-N,N1-bis(aminocarboxylic acid) dialkyl ester with AcCl or AcBr in the presence of a tertiary organic base. Thus, a mixture of equal vols. mucic acid di-Et ester (m. 172°) and H2NCH2CO2Et were left 2 days at room temperature, and the reaction product was washed with ether to give mucyl-N, N1-bis(glycine) diethyl ester (I), m. 205-6° (H2O). A suspension of I 3.8 in pyridine 4 and CHCl3 50 treated at 0-10° with AcCl 3.3 and CHCl3 4 parts, heated 1.5 hrs. to reflux, washed with N HCl, NaHCO3 solution and H2O, dried and concentrated gave tetra-O-acetylmucyl-N,N1-bis(glycine) diethyl ester, m. 216-18° (MeOH). Similar RCO(CHOAc)4COR prepared were (R and m.p. given): l-EtCO2CH2CH2CH(CO2Et)NH, 147-8° (MeOH); l-EtCO2CH2CH(CO2Et)NH, 140-1° (EtOAc); l-MeSCH2CH2CH(CO2Et)NH, 191-2° (EtOAc); EtCO2CH2NMe, 78-9° (alc); l-EtCO2CHMeNH, 192-5° (alc.).
IT Aspartic acid, N,N'-galactaroyldi-, tetraethyl ester, tetraacetate, (-)-
Glutamic acid, N,N'-galactaroyldi-, tetraethyl ester, tetraacetate, L-
Glycine, N,N'-galactoaroyldi-, diethyl ester
Glycine, N,N'-galactoaroyldi-, diethyl ester, tetraacetate
Methionine, N,N'-galactaroyldi-, diethyl ester, tetraacetate, l-
IT 526-99-8, Galactaric acid
(derivs.)
IT 22634-92-0, Heptanedioic acid, 4-oxo-, dimethyl ester 101502-37-8,
Alanine, N,N'-galactaroyldi-, diethyl ester, tetraacetate, L-

101502-38-9, Sarcosine, N,N'-galactaroyldi-, diethyl ester, tetraacetate
(preparation of)

L18 ANSWER 7 OF 7 HCAPLUS COPYRIGHT 2005 ACS on STN
AN 1959:67464 HCAPLUS
DN 53:67464
OREF 53:12185i,12186a-c
ED Entered STN: 22 Apr 2001
TI Tetraacetylmucyl-N,N1-bis(aminomonocarboxylic acid) dialkyl esters and tetraacetylmucyl-N,N1-bis(aminodicarboxylic acid) tetraalkyl esters
PA J. R. Geigy Akt.-Ges.
DT Patent
LA Unavailable
CC 10B (Organic Chemistry: Aliphatic Compounds)
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI GB 807601		19590121	GB	

CLASS

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES

AB The title compds., having antiphlogistic action and useful for the treatment of inflammatory disorders, were prepared H2NCH2CO2Et (I).HCl 14 added to tetraacetylmucic acid dichloride 21 in CHCl3 500, Et3N 22 in CHCl3 75 parts added dropwise with stirring at 0-5°, the mixture stirred 3 hrs. at room temperature, the CHCl3 solution washed with dilute HCl solution, NaHCO3 solution, and H2O, dried, the CHCl3 distilled, and the residue recrystd. from MeOH gave ROC[CH(OAc)]4COR (II) (R = NHCH2CO2Et) (III), m. 216-18°. An alternative procedure for preparing III was described wherein di-Et mucate and I were condensed and the product (m. 205-6°) acetylated. Similarly were prepared the following II (R and m.p. given): L-EtO2CCH2CH2CH(CO2Et)NH, 147-8° (MeOH); L-EtO2CCH2CH(CO2Et)NH, 140-1° (EtOAc); L-MeSCH2CH2CH(CO2Et)NH, 191-2° (EtOAc); EtO2CCH2NMe, 78-9° (EtOH); L-EtO2CCH(Me)NH, 192-5° (EtOH).
IT Inflammation
(-inhibiting substances, N,N-carboxyalkylmucamides as)
IT Alanine, N,N'-mucoyldi-, L-, di-Et ester, tetraacetate
Aspartic acid, N,N'-mucoyldi-, L-, tetra-Et ester, tetraacetate
Glutamic acid, N,N'-mucoyldi-, L-, tetra-Et ester, tetraacetate
Glycine, N,N'-mucoyldi-, diethyl ester, tetraactate
Methionine, N,N'-mucoyldi-, L-, di-Et ester, tetraacetate
IT 101502-38-9, Sarcosine, N,N'-mucoyldi-, diethyl ester, tetraacetate
(preparation of)
IT 5627-20-3, Mucamide
(N,N'-carboxyalkyl derivs.)

=> b hcao

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FILE LAST UPDATED: 01 May 1997 (19970501/UP)

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=> d all l16 tot

L16 ANSWER 1 OF 2 HCAOLD COPYRIGHT 2005 ACS on STN
AN CA56:11449g CAOLD
TI mucic acid derivs.
AU Morel, Charles
PA Geigy, J. R., A.-G.
DT Patent

PATENT NO.	KIND	DATE
CH 354788		

IT 29214-62-8 98075-57-1 101502-37-8 101502-38-9 101502-46-9 106217-01-0 106684-43-9 107541-49-1

L16 ANSWER 2 OF 2 HCAOLD COPYRIGHT 2005 ACS on STN
AN CA53:12186a CAOLD
TI tetraacetylmucyl-N,N'-bis(aminomonocarboxylic acid) dialkyl esters and tetraacetylmucyl-N,N'-bis(aminodicarboxylic acid) tetraalkyl esters
PA Geigy, J. R., A.-G.
DT Patent

PATENT NO.	KIND	DATE
GB 807601		

IT 101502-37-8 101502-38-9 101502-46-9 106217-01-0 106684-43-9 107541-49-1

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